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(54) Title: NOVEL β -ACTIN AND RPS21 PROMOTERS AND USES THEREOF

(57) Abstract: The invention relates to isolation of novel β -actin and ribosomal protein S21 (rpS21) promoters and uses thereof. In particular, this invention features nucleotide sequences for rodent β -actin promoters including, hamster, rat, and mouse, and hamster rpS21 promoter.



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NOVEL β -ACTIN and RPS21 PROMOTERS AND USES THEREOF

[0001] This application claims the benefit of priority of U.S. Provisional Patent Application No. 60/480,768, filed June 24, 2003, the entire contents of which are incorporated by reference.

Field of the Invention

[0002] This invention relates to regulatory gene elements such as promoters and uses thereof, for example, for expression of proteins. More specifically, this invention relates to β -actin and ribosomal protein S21 gene promoters.

Background of the Invention

[0003] Every eukaryotic gene contains regulatory elements driving transcription of that gene. Such regulatory elements include promoters, which are typically positioned immediately upstream of the coding sequence in a gene. Promoters regulate transcription by providing binding sites for transcription factors, which are a part of the transcription machinery. Promoters are commonly used to express proteins in cell culture and in vivo. Many promoters are known and used for expression of proteins in various expression systems. Examples of promoters include cytomegalovirus (CMV) immediate early promoter, Rous sarcoma virus genome large genome long terminal repeats (RSV), Simian Virus 40 (SV40) promoter, interferon gene

promoter, metallothionein promoter, and the thymidine kinase promoter and others, e.g., as described in Fernandez et al. (1999) Gene Expression Systems, Academic Press. However, there is still a need in the art to provide promoters that are capable of generating high levels of expression and/or sustain expression for an extended period of time.

[0004] β -actin is a structural protein and is usually expressed in all species, from protozoa to eukaryotes, including humans. The human and chicken β -actin promoters have been previously described. The β -actin promoter, in general, shows a more ubiquitous activity than the CMV promoter which is widely used (Xu et al. (2001) Gene 272:149-156). The chicken β -actin promoter was shown to exhibit a higher activity than viral CMV and SV40 promoters but only when it is linked to a CMV enhancer sequence (Xu et al., supra).

[0005] The ribosomal protein S21 (rpS21) which is associated with the 40S subunit of the ribosome. The promoter of the human rpS21 gene was previously identified (GenBank® accession No. AJ250907). Similarly to most ribosomal gene promoters, it lacks conventional transcription elements such as the TATA box and CAAT sequence (Smirnova et al. (2000) Bioorg. Khim. 26(5):392-396).

SUMMARY OF THE INVENTION

[0006] This invention provides novel β -actin promoters that have a low level of sequence homology to previously known β -actin promoters (such

as, e.g., human and chicken). This invention further provides novel rpS21 promoters that have a low level of sequence homology to previously known rpS21 promoters (such as, e.g., human and mouse).

[0007] The present invention is based, in part, on the discovery and isolation of β -actin and rpS21 promoters from a Chinese hamster ovary (CHO) cell line. This invention is further based, in part, on an observation that the hamster β -actin promoter has a significantly higher activity than the CMV promoter. The invention is further based, in part, on an observation that the rpS21 promoter is at least as active as the hamster β -actin promoter when used for expressing certain genes. The invention provides nucleotide sequences for these promoters and includes variants of the nucleotide sequences having promoter activity. In some embodiments, a β -actin promoter of the invention is derived from a rodent, for example, hamster, rat, and mouse. The rpS21 promoter is typically derived from a hamster.

[0008] The invention further provides vectors comprising a β -actin or a rpS21 promoter of the invention operably linked to a heterologous nucleic acid. In certain embodiments, a vector of the invention comprises a promoter that is operably linked to a heterologous nucleic acid which encodes a heterologous expression product such as, e.g., a therapeutic protein or a fragment thereof. In illustrative embodiments, the expression product is acid sphingomyelinase (ASM), α -glucosidase (GAA), or tissue plasminogen activator (tPA).

[0009] The invention also provides host cells transfected with a vector of the invention. In illustrative embodiments, the host cell is a mammalian cell such as, e.g., CHO, HEK, and BHK.

[0010] Methods for producing a protein are also provided. Methods for producing a protein include, for example, culturing a cell transfected with a vector comprising a β -actin promoter and/or a rpS21 promoter of the invention operably linked to a heterologous nucleic acid encoding a protein, and recovering the protein. In some embodiments, the heterologous expression product is a secretory protein, which is recovered from the medium. In illustrative embodiments, the protein is ASM, GAA, or tPA.

BRIEF DESCRIPTION OF THE FIGURES

[0011] **Figure 1A** shows an alignment between portions of nucleotide sequences of a hamster β -actin promoter (SEQ ID NO:1) and a rat β -actin promoter (SEQ ID NO:2), demonstrating a 79% identity between nucleotide (nt) 487 to nt 893 of SEQ ID NO:1 and nt 1 to nt 417 of SEQ ID NO:2. The rat β -actin promoter (SEQ ID NO:2) has a 67% identity over the entire length of hamster β -actin promoter (SEQ ID NO:1).

[0012] **Figure 1B** shows an alignment between portions of nucleotide sequences of a hamster β -actin promoter (SEQ ID NO:1) and a rat β -actin promoter (SEQ ID NO:2), demonstrating an 83% identity between nt 1047 to nt 3006 of SEQ ID NO:1 and nt 546 to nt 2493 of SEQ ID NO:2.

[0013] **Figure 2A** shows an alignment between portions of nucleotide sequences of a hamster β -actin promoter (SEQ ID NO:1) and a mouse β -actin promoter (SEQ ID NO:3), demonstrating an 84% identity between nt 33 to nt 487 of SEQ ID NO:1 and nt 1 to nt 449 of SEQ ID NO:3. The mouse β -actin promoter sequence (SEQ ID NO:3) has an 80% identity over the entire length of hamster β -actin promoter sequence of SEQ ID NO:1.

[0014] **Figure 2B** shows an alignment between portions of nucleotide sequences of a hamster β -actin promoter (SEQ ID NO:1) and a mouse β -actin promoter (SEQ ID NO:3), demonstrating an 83% identity between nt 996 to nt 3006 of SEQ ID NO:1 and nt 921 to nt 2953 of SEQ ID NO:1.

[0015] **Figure 3** shows an alignment between portions of nucleotide sequences of a hamster β -actin promoter (SEQ ID NO:1) and a hamster β -actin gene (Genbank® Accession No. U20114; SEQ ID NO:4), demonstrating a 98% identity between nt 1775 to nt 3006 of SEQ ID NO:1 and nt 1 to nt 1232 of SEQ ID NO:4. The hamster β -actin gene sequence has a 40% identity over the entire length of the hamster β -actin promoter sequence of SEQ ID NO:1.

[0016] **Figure 4** shows an alignment between portions of nucleotide sequences of hamster β -actin promoter (SEQ ID NO:1) and a previously known human β -actin promoter (GenBank® Accession No. gi28337; SEQ ID NO:5), demonstrating a 94% identity between nt 113 to nt 148 of SEQ ID NO:1 and nt 38 to nt 73 of SEQ ID NO:5, an 83% identity

between nt 362 to nt 433 of SEQ ID NO:1 and nt 303 to nt 374 of SEQ ID NO:5, a 90% identity between nt 1728 to nt 1764 of SEQ ID NO:1 and nt 1791 and nt 1830 of SEQ ID NO:5, and a 91% identity between nt 1797 to nt 1966 of SEQ ID NO:1 and nt 1840 to nt 2007 of SEQ ID NO:5. The human β -actin promoter sequence (SEQ ID NO:5) shows a 10% identity over the entire length of the hamster β -actin promoter sequence of SEQ ID NO:1.

[0017] **Figure 5** shows an alignment between portions of nucleotide sequences of hamster β -actin promoter (SEQ ID NO:1) and a previously known chicken β -actin promoter (GenBank® Accession No. gi2170437; SEQ ID NO:6), demonstrating an 83% identity between nt 1878 to nt 1919 of SEQ ID NO:1 and nt 186 to nt 227 of SEQ ID NO:6. The chicken β -actin promoter sequence (SEQ ID NO:6) shows a 1% identity over the entire length of the hamster β -actin promoter sequence of SEQ ID NO:1.

[0018] **Figure 6A** depicts a Northern blot for galectin, ferritin, and β -actin in CHO-K1 cells. Representative mRNAs were isolated from cells at 0, 4, 8, 10, and 15 hours following treatment of cells with actinomycin D.

[0019] **Figure 6B** depicts relative mRNA expression levels for galectin, ferritin, and β -actin genes. Representative mRNAs were isolated from cells at 0, 4, 8, 10, and 15 hours following treatment of CHO-K1 cells with actinomycin D.

[0020] **Figure 7A** depicts relative promoter strengths as measured in transient transfection assays in CHO-K1 cells for the following promoters: CMV, human EF-1, hamster GAPDH, hamster rpS21 and hamster β -actin.

The representative promoters were cloned upstream of a red fluorescent protein (RFP) gene in the pDsRED-1 plasmid. The mean fluorescence was measured by FACS.

[0021] **Figure 7B** depicts relative promoter strengths as measured in stable transfection assays in CHO-K1 cells for the following promoters: CMV, human EF-1, hamster GAPDH, hamster rpS21, and hamster β -actin. The representative promoters were cloned upstream of a red fluorescent protein (RFP) gene in the pDsRED-1 plasmid. The mean fluorescence was measured by FACS.

[0022] **Figure 8A** depicts the expression of acid sphingomyelinase (ASM) protein in media from three pools of CHO-DXB11 cells transfected with a vector containing the ASM cDNA operably linked to either the CMV promoter or the hamster β -actin promoter. The expression of ASM was assessed in an enzymatic activity assay for ASM.

[0023] **Figure 8B** depicts the expression of α -glucosidase (GAA) protein in media from three pools of CHO-DXB11 cells transfected with a vector containing the GAA cDNA operably linked to either the CMV promoter or the hamster β -actin promoter. The expression of GAA was assessed in an enzyme activity assay for GAA.

[0024] **Figure 9** depicts the expression of tPA protein in media from pools of CHO-DXB11 cells transfected with a vector containing the tPA cDNA operably linked to the hamster β -actin promoter. The expression of tPA was assessed using ELISA.

DETAILED DESCRIPTION OF THE INVENTION

[0025] In order that the present invention be more readily understood, certain terms are first defined. Additional definitions are set forth throughout the detailed description.

[0026] The term "promoter" refers to a regulatory element that directs the transcription of a nucleic acid to which it is operably linked. A promoter can regulate both rate and efficiency of transcription of an operably linked nucleic acid. A promoter may also be operably linked to other regulatory elements which enhance ("enhancers") or repress ("repressors") promoter-dependent transcription of a nucleic acid. The term "operably linked" refers to a nucleic acid placed in a functional relationship with another nucleic acid. A promoter is usually positioned 5' (i.e., upstream) of a transcription initiation site in the nucleic acid. A promoter, however, may include sequences 3' (i.e., downstream) of the transcription initiation site. A promoter may also encompass regions both 5' and 3' of the transcription initiation site of the operably linked nucleic acid.

[0027] The term "promoter activity" refers to the ability of a promoter to initiate transcription of a nucleic acid to which it is operably linked. Promoter activity can be measured using procedures known in the art or as described in the Examples. For example, promoter activity can be measured as an amount of mRNA transcribed by using, for example, Northern blotting or polymerase chain reaction (PCR). Alternatively, promoter activity can be measured as an amount of translated protein product, for example, by

Western blotting, ELISA, colorimetric assays such as, e.g., Bradford assay (Bradford (1976) Anal. Biochem., 72:248), and various activity assays, including reporter gene assays and other procedures known in the art or as described in the Examples.

[0028] The term "vector" refers to viral or non-viral, prokaryotic or eukaryotic, deoxyribonucleic acid, ribonucleic acid or a nucleic acid analog, that is capable of carrying another nucleic acid. A vector may either carry a nucleic acid into a cell, referred to as "host cell," so that all or a part of the nucleic acid is transcribed or expressed. Alternatively, a vector may be used in an in vitro transcription assay. Vectors are frequently assembled as composites of elements derived from different viral, bacterial, or mammalian genes. Vectors contain various coding and non-coding sequences including sequences coding for selectable markers (e.g., an antibiotic resistance gene), sequences that facilitate their propagation in bacteria, or one or more transcription units that are expressed only in certain cell types. For example, mammalian expression vectors often contain both prokaryotic sequences that facilitate the propagation of the vector in bacteria and one or more eukaryotic transcription units that are expressed only in eukaryotic cells. It will be appreciated by those skilled in the art that the design of the expression vector can depend on such factors as the choice of the host cell to be transformed, the level of expression of protein desired, etc.

[0029] Vectors include, for example, plasmids, phagemids, and viral vectors. Vectors that have an existing promoter can be modified by

standard recombinant DNA techniques known in the art to replace the promoter with any of promoter sequences set forth in SEQ ID NOs:1, 2, 3, or 39 or a variant thereof. In general, suitable vectors can either be chosen from those that are commercially available or they can be constructed using standard recombinant DNA techniques known in the art. (See, e.g., Molecular Cloning: A Laboratory Manual: 2nd edition, Sambrook et al., 1989, Cold Spring Harbor Laboratory Press.)

[0030] The terms "transformation" and "transfection" refer to intracellular introduction of a nucleic acid. A nucleic acid can be introduced into a plant or an animal cell or a prokaryotic or eukaryotic cell by a number of methods known in the art or described herein.

[0031] The term "isolated" refers to a deoxyribonucleic acid, a ribonucleic acid, or a nucleic acid analog having a polynucleotide sequence that is separated from other nucleic acid sequences in such a way that does not naturally occur. An isolated nucleic acid encompasses nucleic acids that may be partially or wholly chemically or recombinantly synthesized and/or purified by standard techniques known in the art.

[0032] The term "variant" in reference to a promoter sequence refers to a nucleotide sequence that is substantially identical over the entire length to the promoter sequence or to its complementary strand over the entire length thereof, provided that the variant has promoter activity.

[0033] Variants of β -actin promoters may be the same length as the nucleotide sequences of SEQ ID NOs:1, 2, or 3, or shorter, so long as

they are at least 1250 nucleotides in length. Variants of rpS21 promoters may be the same length as the nucleotide sequence of SEQ ID NO:39, or shorter, so long as they have promoter activity. Variants of the β -actin promoter can be naturally occurring, for example, naturally occurring β -actin promoters isolated from species other than human and chicken, or they can be generated artificially. The identity between the hamster β -actin promoter set forth in SEQ ID NO:1 and a variant thereof, when optimally aligned, is at least 45%, 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% over the entire sequence of SEQ ID NO:1 from nt 1 to nt 3007. Similarly, the identity between the rat β -actin promoter set forth in SEQ ID NO:2 and a variant thereof is at least 60%, 65%, 70%, 75%, 80%, 85%, 90%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% over the entire sequence of SEQ ID NO:2 from nt 1 to nt 2493. The identity between the mouse β -actin promoter of SEQ ID NO:3 and a variant thereof is at least 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% over the entire length of SEQ ID NO:3 from nt 1 to nt 2953. Similarly, identity between the hamster rpS21 promoter set forth in SEQ ID NO:39 and a variant thereof, when optimally aligned, can be at least 40%, 50%, 55%, 60%, 65%, 70%, 80%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% over the entire length of SEQ ID NO:39 from nt 1 to nt 1958.

[0034] Variants of β -actin promoters may, for example, include orthologs of the β -actin promoters in other species, including rodents and

other mammals, but excluding human and chicken β -actin promoters and known variants thereof. Variants of the promoters of the invention may also be found in other rodent species such as, for example, guinea pig, woodchuck, muskrat, gerbil, squirrel, chipmunk, prairie dog, beaver, porcupine, and vole.

[0035] The term "variants" further encompasses fragments of any one or more of promoters of the invention that have promoter activity. Variants of the β -actin promoters are at least 1250 nucleotides in length. Variants of the β -actin promoters of the invention can be derived, for example, by 5' truncations of the hamster β -actin promoter set forth in SEQ ID NO:1. In some embodiments, β -actin promoter variants include sequences from nt 50 to nt 3000, from nt 100 to nt 3000, from nt 150 to nt 3000, from nt 200 to nt 3000, from nt 250 to nt 3000, from nt 500 to nt 3000, from nt 1000 to nt 3000, or from nt 1500 to nt 3000 of SEQ ID NO:1. In other embodiments, β -actin promoter variants may be derived by 5' truncations of the sequence set forth in SEQ ID NO:2 and include, for example, from nt 50 to nt 2490, from nt 100 to nt 2490, from nt 150 to nt 2490, from nt 200 to nt 2490, from nt 250 to nt 2490, from nt 500 to nt 2490, or from nt 1000 to nt 2490 of SEQ ID NO:2. β -actin promoter variants may also be derived by 5' truncations of the sequence set forth in SEQ ID NO:3 and include, for example, from nt 50 to nt 2950, from nt 100 to nt 2950, from nt 150 to nt 2950, from nt 200 to nt 2950, from nt 250 to nt 2950, from nt 500 to nt 2950, from nt 1000 to nt 2950, or from nt 1500 to nt 2950 of SEQ ID NO:3. Longer fragments of the hamster

β -actin promoter can be derived, for example, by 5' truncations of the longer hamster promoter nucleotide sequence set forth in SEQ ID NO:7. Such variants include, for example, sequences from nt 50 to nt 3668, from nt 100 to nt 3668, from nt 150 to nt 3668, from nt 200 to nt 3668, from nt 250 to nt 3668, from nt 500 to nt 3668, or from nt 600 to nt 3668.

[0036] Variants of rpS21 promoters may be derived by 5' truncations and/or 3' truncations of the sequence set forth in SEQ ID NO:39. Such variants include, for example, sequences from nt 50 to nt 1958, from nt 100 to nt 1958, from nt 150 to nt 1958, from nt 200 to nt 1958, from nt 250 to nt 1958, from nt 500 to nt 1958, from nt 1000 to nt 1958, from nt 1 to nt 1900, from nt 1 to nt 1850, from nt 1 to nt 1800, from nt 1 to nt 1750, from nt 1 to 1700, from nt 1 to nt 1600, or from nt 1 to nt 1500.

[0037] In certain embodiments, a β -actin promoter of the invention comprises a contiguous stretch of at least 1250, 1500, 1550, 1600, 1650, 1700, 1750, 1800, 1850, 1900, 1950, 2000, 2500, or 3000 nucleotides from SEQ ID NOs:1, 2, or 3. Such contiguous stretches of SEQ ID NOs:1, 2, and 3 may also contain a mutation (insertion or deletion) so long as the mutant sequence retains at least some functionality of the original sequence and the capacity to hybridize to the respective sequences of SEQ ID NOs:1, 2, or 3 under low, medium or high stringency conditions. A contiguous stretch of a β -actin promoter can be derived by 5' truncations of any of sequences set forth in SEQ ID NO:1, 2, 3, or 7 or variants thereof as described above.

[0038] In other embodiments, a rpS21 promoter of the invention comprises a contiguous stretch of at least 500, 600, 700, 800, 900, 1000, 1100, 1200, 1300, 1400, 1500, 1600, 1700, 1800, 1850, or 1900 nucleotides from SEQ ID NO:39.

[0039] β -actin promoter variants of the invention further include nucleotide sequences that hybridize to the entire length of the β -actin promoter sequences shown in SEQ ID NOs:1, 2, or 3, or their complements and that have at most 0, 1, 2, 3, 4, 5, 10, 15, 20, 25, 30, 35, 40, 45% base pair mismatches. rpS21 promoter variants of the invention include nucleotide sequences that hybridize to the entire length of the rpS21 promoter sequence shown in SEQ ID NO:39, or its complement, and that have at most 0, 1, 2, 3, 4, 5, 10, 15, 20, 30, 40, 45, 50, 55, 60% base pair mismatches. The percentage of base pair mismatches can be determined by standard techniques known in the art or as described herein. The term "heterologous" when used in reference to a nucleic acid, means a nucleic acid other than the nucleic acid that a promoter is operably linked to in a naturally occurring genome. For example, the term "heterologous" refers to any nucleic acid other than the hamster β -actin gene when such a nucleic acid is operably linked to a hamster β -actin promoter. Likewise, the term "heterologous" refers to any nucleic acid other than the rat β -actin gene when such a nucleic acid is operably linked to a rat β -actin promoter. Similarly, the term "heterologous" refers to any nucleic acid when such a nucleic acid is operably linked to the mouse β -actin promoter. Analogously, this term also refers to any nucleic

acid other than the hamster rpS21 gene when such a nucleic acid is operably linked to a hamster rpS21 promoter.

[0040] The term "transgenic" refers to any animal containing genetically manipulated cells in which a promoter of the invention is no longer operably linked to the same nucleic acid as in a naturally occurring genome. The term "transgenic" encompasses, for example, an animal containing cells with a promoter of the invention or a variant thereof integrated within the animal's chromosome. The term "transgenic" also encompasses an animal containing cells with an extrachromosomally replicating DNA sequence comprising a promoter of the invention or a variant thereof. The transgenic animal may be a mammal such as a rodent or human.

[0041] This invention is based, in part, on the discovery and isolation of novel promoters for the β -actin and rpS21 genes. Specifically, this invention features rodent β -actin promoters including, but not limited to, hamster, rat and mouse, and the hamster rpS21 promoter. This invention is based on the discovery and demonstration that β -actin promoters of the invention have promoter activity that is higher than the CMV promoter's activity, as described in the Examples. The invention is further based on the discovery that the hamster rpS21 promoter is at least as active as the hamster β -actin promoter when used for expressing certain genes.

[0042] The invention provides nucleotide sequences for rodent β -actin promoters, including hamster, rat, and mouse, and methods of use thereof. The invention further provides methods for identification and isolation

of variants of promoters of the invention, including homologs and fragments of promoters that have promoter activity. Additionally, the invention provides a nucleotide sequence for the hamster rpS21 promoter, and methods of use thereof.

[0043] In the experiments leading to the present invention, a genomic clone for the hamster β -actin promoter was isolated from CHO cells following its identification as an active promoter by a technique called Serial Analysis of Gene Expression or "SAGE" (Valculesco et al. (1995) Science, 270:484-487 and Valculesco et al. (1987) Cell, 88:243-251). The SAGE technique can be used for transcription profiling of an entire genome. β -actin promoter was identified as one of the most active promoters in CHO cells using SAGE. This led to the cloning of the promoter for β -actin in CHO cells. A similar approach was used for the isolation of the hamster rpS21 promoter from CHO cells. This approach may be used for transcription profiling of other genomes to confirm that corresponding β -actin promoters or rpS21 promoter are active in another genome. Such a promoter can be cloned using standard techniques known in the art or those described here. Variants of promoters of the invention can be identified by hybridization to one or more of promoter sequences set forth in SEQ ID NOs:1, 2, 3, or 39. It is well known that the melting temperature (T_m) of a double-stranded nucleic acid decreases by 1-1.5°C with every 1% decrease in homology (see, e.g., Bonner et al. (1973) J. Mol. Biol., 81:123). Species homologs, therefore, can be identified, for example, by hybridizing a putative nucleotide sequence with a nucleotide

sequence of SEQ ID NOs:1, 2, 3, or 39, or a variant thereof, and comparing the melting temperature of such a hybrid with the melting temperature of a hybrid comprising a nucleotide sequence of SEQ ID NOs:1, 2, 3, or 39, or a variant thereof and a complementary nucleotide sequence. The number of base pair mismatches can then be calculated for the test hybrid. Therefore, a smaller difference between the melting temperatures of the test hybrid and a hybrid containing a putative homolog of any one of sequences in SEQ ID NOs:1, 2, 3, or 39, will indicate a greater homology between the putative nucleotide sequence and a promoter sequence of the invention. For example, variants in other rodent species such as guinea pig, woodchuck, muskrat, gerbil, squirrel, chipmunk, prairie dog, beaver, porcupine, and vole, may exhibit a greater homology to promoters of the invention and variants thereof.

[0044] A variety of factors are known to affect the efficiency of hybridization of two strands of nucleotide sequence. These may include, for example, length of nucleotide sequence, salt concentration and G/C content of the sequences. For example, for hybridization of long fragments of DNA, Howley et al. (1979) J. Biol. Chem., 254:4876, determined that the melting temperature at which 50% of a DNA is hybridized to a complementary strand is defined by:

$$T_m = 81.5 + 16.6 \log M + 41(\%G + \%C) - 500/L - 0.62F,$$

where

M is molar concentration of monovalent cations;

(%G + %C) is the respective fraction of G and C nucleotides in the sequences;

L is length of the hybrid DNA; and

F is molar concentration of formamide.

[0045] Appropriate hybridization conditions can be selected by those skilled in the art with minimal experimentation as exemplified in Ausubel et al. (1995) Current Protocols in Molecular Biology, John Wiley & Sons, sections 2, 4, and 6. Additionally, stringent conditions are described in Sambrook et al. (1989) Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor Press, chapters 7, 9, and 11.

[0046] A non-limiting example of low stringency hybridization conditions is as follows. Filters containing DNA are pretreated for 6 h at 40°C. in a solution containing 35% formamide, 5 × SSC, 50 mM Tris-HCl (pH 7.5), 5 mM EDTA, 0.1% PVP, 0.1% Ficoll™, 1% BSA, and 500 µg/ml denatured salmon sperm DNA. Hybridizations are carried out in the same solution with the following modifications: 0.02% PVP, 0.02% Ficoll™, 0.2% BSA, 100 µg/ml salmon sperm DNA, 10% (wt/vol) dextran sulfate, and 5-20 × 10⁶ 32P-labeled probe is used. Filters are incubated in hybridization mixture for 18-20 h at 40°C, and then washed for 1.5 hours at 55°C in a solution containing 2 × SSC, 25 mM Tris-HCl (pH 7.4), 5 mM EDTA, and 0.1% SDS. The wash solution is replaced with fresh solution and incubated for an additional 1.5 hours at 60°C. Filters are blotted dry and exposed for autoradiography. Other conditions of

low stringency well known in the art may be used (e.g., as employed for cross species hybridizations).

[0047] A non-limiting example of high stringency hybridization conditions is as follows. Prehybridization of filters containing DNA is carried out for 8 h to overnight at 65°C in buffer containing 6 × SSC, 50 mM Tris-HCl (pH 7.5), 1 mM EDTA, 0.02% PVP, 0.02% Ficoll™, 0.02% BSA, and 500 µg/ml denatured salmon sperm DNA. Filters are hybridized for 48 hours at 65°C in the prehybridization mixture containing 100 µg /ml denatured salmon sperm DNA and 5-20 × 10⁶ cpm of ³²P-labeled probe. Washing of filters is done at 37°C for 1 hours in a solution containing 2 × SSC, 0.01% PVP, 0.01% Ficoll™, and 0.01% BSA. This is followed by a wash in 0.1 × SSC at 50°C for 45 minutes.

[0048] A non-limiting example of hybridization conditions of moderate stringency includes prewashing filters in 5 × SSC, 0.5% SDS, 1.0 mM EDTA, pH 8.0; hybridizing in 50% formamide, 6 × SSC at 42°C; and washing filters in 0.5 × SSC, 0.1% SDS at 60°C.

[0049] Variants of the promoters of the invention can also be identified by percent identity between nucleotide sequences for putative variants and the sequences set forth in SEQ ID NOs:1, 2, 3, or 39, or their complementary strands. Percent identity may be determined, for example, by visual inspection or by using various computer programs known in the art or as described in the Examples. For example, percent identity of two nucleotide sequences can be determined by comparing sequence information using the

GAP computer program described by Devereux et al. (1984) Nucl. Acids. Res., 12:387 and available from the University of Wisconsin Genetics Computer Group (UWGCG). Percent identity can also be determined by aligning two nucleotide sequences using the BLAST® program (www.ncbi.nlm.nih.gov/BLAST) as described by Tatusova et al. (1999) FEMS Microbiol. Lett., 174:247. For example, for nucleotide sequence alignments using the BLAST® program, the default settings are as follows: reward for match is 2, penalty for mismatch is -2, open gap and extension gap penalties are 5 and 2 respectively, gap × dropoff is 50, expect is 10, word size is 11, and filter is OFF.

[0050] Promoters of the invention identified by sequence identity include, for example, sequences set forth in SEQ ID NOs:2 and 3 for rat and mouse β -actin promoters, that show 67% and 80% identity, respectively, to nt 1 to nt 3007 of hamster β -actin promoter sequence set forth in SEQ ID NO:1. Additional variants can be readily identified using the various techniques described herein and those known in the art.

[0051] Percent identity between the hamster β -actin promoter (SEQ ID NO:1) and known β -actin promoters can be determined as described. For example, when SEQ ID NO:1 is compared to the human β -actin promoter (SEQ ID NO:5) using BLAST® sequence alignment with default parameters, it exhibits only about a 10% identity over the entire length of SEQ ID NO:1. Similarly, when SEQ ID NO:1 is compared to the chicken β -actin promoter (SEQ ID NO:6), it exhibits only about a 1% identity over the

entire length of SEQ ID NO:1. Due to such low levels of homology, the human and the chicken β -actin promoters are not considered to be variants of the hamster β -actin promoter sequence of SEQ ID NO:1. Further, the 3' portion of SEQ ID NO:1 shows significant homology to the 5' portion of the hamster β -actin gene sequence (GenBank® Accession No. U20114; SEQ ID NO:4). In particular, the first 1232 nucleotides of SEQ ID NO:4 show a 98% identity with the 3' portion of SEQ ID NO:1, as depicted in Figure 3. This identity is in the region of the first intron in the hamster β -actin gene. Overall, SEQ ID NO:4 shows only 40% identity over the entire length of SEQ ID NO:1. Furthermore, no promoter activity has been described for SEQ ID NO:4, or fragments thereof.

[0052] Using BLAST® sequence alignment with default parameters, no homology is detected between the previously known human rpS21 promoter (nt 1-2344 of GenBank® Accession No. AJ250907) and nt 1 to 1958 of hamster rpS21 promoter of SEQ ID NO:39. Very low level of homology is detected between hamster rpS21 promoter of SEQ ID NO:39 and mouse genomic DNA that spans the mouse rpS21 gene (GenBank® Accession No. NT_039212). There are two regions of homology in the mouse sequences. The first is from nt 1775 to nt 1945 of SEQ ID NO:39 (137 out of 172 nts match). The second is from nt 580 to nt 851 of SEQ ID NO:39 (208 out of 274 nts match). These two regions of homology are separated by 923 nts in the hamster sequence (SEQ ID NO:39) and by 1745 nts in the mouse genomic sequence (NT_039212).

[0053] Accordingly, in some embodiments, an isolated promoter or a variant thereof having promoter activity comprises the nucleotides sequence(s) as set out from nt 1775 to nt 1945 of SEQ ID NO:39 and/or from nt 580 to nt 851 of SEQ ID NO:39. Optionally, such a promoter or variant further comprises all or a portion of SEQ ID NO:39 as set out from nt 852 to nt 1774.

[0054] Nucleotide sequences set forth in SEQ ID NOs:1, 2, 3, or 39, or variants thereof, can be used as probes for screening genomic libraries for the isolation of genomic sequences that hybridize to one or more of sequences set forth in SEQ ID NOs:1, 2, 3, or 39, or variants thereof.

[0055] A promoter, according to the invention, or a variant thereof is operably linked to a heterologous nucleic acid which it expresses. The promoter can be used either alone or in combination with other regulatory elements such as, for example, enhancers and repressors. Alternatively, such a promoter can be integrated into the genome of a host cell or animal, thereby to express an endogenous gene in the host. A promoter according to the invention can be used in a vector for expression of heterologous nucleic acids. In certain embodiments, the heterologous nucleic acid encodes a therapeutic protein. Examples of therapeutic proteins include, but are not limited to, α -glucosidase, acid sphingomyelinase, insulin, tissue plasminogen activator, thyrogen stimulating hormone, erythropoietin, glucocerebrosidase, α -galactosidase and various antibodies. Examples of antibodies include but

are not limited to, antibodies that bind members of the TGF- β family such as, for example, TGF- β -1, 2, and 3.

[0056] This invention further provides vectors comprising a promoter of the invention or a variant thereof which has promoter activity. In some embodiments, vectors of the invention include a suitable restriction enzyme site downstream of the promoter for insertion of the heterologous nucleic acid. Such a restriction enzyme site may include a restriction site for a single restriction enzyme or it may include restriction sites for a variety of restriction enzymes in order to facilitate insertion of many different heterologous nucleic acids. A vector according to the invention may also contain a polyadenylation sequence downstream of the site for inserting a heterologous nucleic acid. Vectors comprising promoters of the invention may also contain prokaryotic DNA elements for bacterial replication and an antibiotic selection marker for growth and selection of the vector in bacterial cells and additional DNA elements that control processing of transcripts such, e.g., termination signals. Vectors may further contain DNA sequences to direct secretion of a protein outside host cells.

[0057] In certain embodiments, a vector containing a promoter sequence of the invention is a bicistronic vector. Bicistronic vectors are designed, such that two nucleic acids can be transcribed to yield a single transcript. Such a transcript usually contains a first portion which is translated into one protein and a second portion translated into a second protein. One protein can be a protein of interest such as, a therapeutic protein, and a

second protein may be used as a selectable marker. Bicistronic vectors usually contain a promoter and an internal ribosome entry site or IRES positioned between two nucleic acids. This permits transcription of the two nucleic acids as a single bicistronic mRNA. In this manner, a vector can be constructed that includes a β -actin promoter of the invention or a variant thereof and an IRES between two heterologous nucleic acids. A bicistronic vector containing a β -actin promoter of the invention or a variant thereof can be used for expressing a therapeutic protein such as, for example, acid sphingomyelinase or α -glucosidase, in conjunction with a reporter gene.

[0058] The invention further provides assays for identification of those variants of β -actin and rpS21 promoters of the invention that have promoter activity. For example, a promoter of the invention or variant thereof is inserted in a suitable vector upstream of a reporter gene and the expression of the reporter gene is used as a determinant of promoter activity. For example, for identification of variants of promoters of the invention that have promoter activity, such a variant is cloned upstream of a reporter gene. A reporter gene may encode an enzyme which catalyzes a reaction which produces a visually detectable signal. Examples of such reporter genes include β -galactosidase and luciferase. Examples of other reporter genes include alkaline phosphatase, nopaline synthase, octopine synthase, β -glucuronidase, chloremphenicol acetyltransferase. In the Examples set forth below, a reporter gene encoding a *Discosoma striata* red fluorescent protein (RFP) is used for measuring promoter activity. Those skilled in the art,

however, can use any suitable reporter gene and assay technique to determine promoter activity. Expression of a reporter gene from the promoter may be assayed in an in vitro expression system or it may be intracellular (e.g., in vivo).

[0059] The invention further provides host cells that have been transfected with a vector of the invention comprising a promoter operably linked to a heterologous gene. Such a host cell can be a prokaryotic cell or a eukaryotic cell. Host cells can either be cells in culture or be present in an animal. Examples of host cells in culture include, but are not limited to, HeLa cells, CHO cells, NS0, HEK cells, BHK cells, NIH-3T3, MDCK cells, and COS cells. Host cells in culture can be grown either in suspension or on microcarriers, as described in the Examples.

[0060] Many suitable methods can be used for introducing nucleic acids of the invention into a host cell. Vectors comprising promoter sequences of the invention can be introduced into either prokaryotic or eukaryotic cells. Examples of techniques that may be used for introduction of nucleic acids into eukaryotic cells include, for example, calcium phosphate precipitation, DEAE-Dextran transfection, electroporation, liposome-mediated transfection, transduction using viral vectors, etc.

[0061] Many suitable expression systems can be employed for the production of proteins using promoters of the invention. One such expression system employs a dihydrofolate reductase (DHFR) gene which is introduced into the vector comprising a promoter of the invention or a variant thereof

operably linked to a heterologous nucleic acid. Alternatively, an expression vector expressing DHFR can be co-transfected into the host cell, if a DHFR-deficient cell is used for expression. When increasing concentrations of methotrexate (MTX), a competitive inhibitor of the essential enzyme DHFR, are applied to the transfected cells, only cells with higher expression levels of DHFR survive. As MTX levels are increased further, only cells which amplify the copy number of the DHFR gene survive. In this way, by increasing the copy number of the vector comprising the promoter, increased expression of the heterologous nucleic acid can be achieved, thereby leading to increased protein production. A second expression system employs a glutamine synthetase (GS) gene that is introduced into the vector comprising a promoter of the invention or a variant thereof operably linked to a heterologous nucleic acid. Addition of a competitive inhibitor of GS, e.g., methionine sulfoximine (MSX), is used for increasing the copy number of the vector leading to increased protein production.

[0062] Any suitable prokaryotic or eukaryotic expression system can be used for expression of proteins using promoters of the invention. Examples of expression systems include, but are not limited to, plant, baculovirus, yeast, bacterial, drosophila, mammalian and cell free expression systems. Standard methods for introducing expression vectors into mammalian, bacterial, yeast, insect and plant cells are provided, for example, by Ausubel (1995), supra.

[0063] In certain embodiments, promoters of the invention and variants thereof are used in methods of gene therapy. For example, a promoter of the invention or a variant thereof is cloned into a viral or a non-viral gene therapy vector such that it is operably linked to a gene of interest. The promoter drives expression of the gene encoding a therapeutic protein when the vector is delivered to a subject, e.g., a human patient.

[0064] The following examples provide illustrative embodiments of the invention. One of ordinary skill in the art will recognize the numerous modifications and variations that may be performed without altering the spirit and scope of the present invention. Such modifications and variations are encompassed within the scope of the invention. The examples do not in any way limit the invention.

EXAMPLES

[0065] The following describes materials and methods used in the subsequent Examples.

A. Culturing of CHO-K1 cells

[0066] CHO-K1 cells were obtained from American Type Culture Collection (Manassas, VA) (ATCC No. CRL-9618). Cells were cultured in 250 ml spinner cultures containing 15 g/L DE-52 microcarriers (Whatman, Kent, UK) in 925 cell culture medium supplemented with 10% donor calf serum (DCS) (Invitrogen). Cells were maintained at 37°C using a 20-40% O₂ and 5% CO₂ overlay and agitated at approximately 60 rpm for six days. Following

growth of cells in the presence of serum, cultures were subjected to a daily 80% (v/v) replacement with serum-free 925 medium. Cells were grown in serum-free medium for 11 days prior to extraction of RNA from cells. For the determination of mRNA half-life, 7 mg/L of actinomycin D was added to the cultures in the serum-free phase.

B. RNA Extraction and Analysis

[0067] RNA was isolated from CHO-K1 cells using the RNAgents kit from Promega (Madison, WI). Gene expression was analyzed by Northern blotting. For Northern blot analysis, 5 µg of RNA was separated by electrophoresis on a denaturing glyoxal/dimethylsulfoxide gel using a NorthernMax®-Gly kit. (Ambion, Austin, TX). The RNA was subsequently transferred to nylon membranes (Schleicher & Schuell, Dassel, Germany). The blots were probed with the following gene probes amplified by PCR: galectin (GenBank® Accession No. M96676, nt 14-383); β-actin (Genbank® Accession No. U20114, nt 238-381); EF-1 (GenBank® Accession No. D00522, nt 7-192); rpS21 (GenBank® Accession No. X79059, nt 68-340); ferritin (GenBank® Accession No. M99692, nt 182-303) or a commercially available glyceraldehyde 3-phosphate dehydrogenase (GAPDH) fragment (Ambion, Austin, TX). Each PCR product was radiolabeled by random priming. PCR primers used for amplification of each of the genes are listed in Table 1.

TABLE 1

Gene	Primer	Sequence	SEQ ID NO:
β -actin	forward	GCTCTTTCTTCGCCGCTCC	8
β -actin	reverse	ACCACCCTCCAGCCTTCCC	9
EF-1	forward	GAACGCAGGTGTTGTGAAAA	10
EF-1	reverse	CTCGGCAGCCTCCTTCT	11
rpS21	forward	GTGGACCTGTACGTGC	12
rpS21	reverse	TTCTCACTTTTATTTATGAC	13
ferritin	forward	CGCCAGAACTACCACCAGGAC	14
ferritin	reverse	TTCAGAGCCACATCATCCCG	15
galectin	forward	TGGT \overline{C} GCAAGCAACCTGAATC	16
galectin	reverse	TTGAAGTCACCGTCTGCCGC	17

C. Transfection of CHO-K1 Cells

[0068] For transient transfection, CHO-K1 cells were plated on 6-well plates in 925 medium with 10% fetal bovine serum (FBS) (Invitrogen). The cells were grown to 50-75% confluency prior to transfection using Lipofectamine™ (Invitrogen). The pDsRED-1 plasmid (Clontech, Palo Alto, CA) was co-transfected with the pSV40-CD20 plasmid, which encodes a cell surface CD20 marker used to identify transfected cells. This pDsRED-1 plasmid encodes a *Discosoma striata* red fluorescent protein (RFP), the expression of which can be detected by FACS. Transfections were performed as per manufacturer's instructions. Briefly, cells were incubated with lipid-DNA complexes for 16 hrs in serum free Opti-MEM™ medium (Invitrogen). The medium was replaced with 925 medium with 10% FBS, and cells were harvested 48 hours post-transfection.

D. Fluorescence-Activated Cell Sorting Analysis

[0069] For FACS analysis, 1×10^6 cells were trypsinized and washed with cold PBS containing 2% FBS. Cells were subsequently incubated with an FITC-labeled anti-CD20 antibody (Pharmingen, San Diego, CA) for 30 minutes on ice. Cells were then washed with cold PBS containing 2% FBS and resuspended in 1 ml of cold PBS/2% FBS. FACS analysis was performed using FACSCalibur™ (BD Biosciences, San Diego, CA). All CD20-positive events were evaluated for their red fluorescent protein mean fluorescence intensity to assess promoter strength.

E. ASM Assay

[0070] Media from cells transfected with a vector encoding acid sphingomyelinase (ASM) were incubated at 37°C with the synthetic substrate 2-(N-hexadecanoylamino)-4-nitrophenylphosphorylchlorine (Calbiochem, San Diego, CA) at the concentration of 12.5 mM in 250 mM sodium acetate, pH 5.5, containing 0.1 mM zinc acetate, 0.25 mg/ml bovine serum albumin (BSA) and 0.15% Tween 20. The reactions were stopped by the addition of 0.2 M glycine-NaOH containing 50% ethanol. The activity or amount of ASM was measured by the amount of 2-(N-hexadecanoylamino)-4-nitrophenolate produced using a colorimetric assay by measuring optical density at 415 nm.

F. GAA Assay

[0071] Media from cells transfected with a vector encoding α -glucosidase (GAA) were incubated at 37°C with the synthetic substrate p-nitrophenyl-D-a-glucopyranoside (Sigma, St. Louis, MO) at a concentration

of 40 mM in 50 mM sodium acetate, pH 4.3, containing 0.1% bovine serum albumin (BSA). The reactions were stopped by the addition of 0.3 M glycine, pH 10.6. The activity or amount of GAA was measured by the amount of p-nitrophenyl produced using a colorimetric assay by measuring optical density at 400 nm.

Example 1: Identification of the β -Actin Promoter in CHO-K1 Cells

[0072] Serial Analysis of Gene Expression (SAGE) was used to analyze the entire transcription profile of CHO-K1 cells that were grown in a serum-free perfused spinner culture.

[0073] The first step in SAGE involved synthesis of double stranded DNA from mRNA isolated from CHO-K1 cells using standard techniques. The cDNA was subsequently cleaved with a restriction endonuclease *NlaIII*, also called an anchoring enzyme, which is expected to cleave most transcripts at least once. The 3' portion of each cleaved cDNA was isolated by binding to streptavidin beads. The cDNA pool was then divided in half and ligated via anchoring the restriction site to a linker containing a type II restriction endonuclease site (for example, *FokI*). Type II restriction endonucleases cleave at a defined distance up to 20 base pairs away from their asymmetric recognition sites. The type II enzyme is typically called a tagging enzyme. Cleavage of the ligation product with the tagging enzyme results in the release of the linker with short pieces of the cDNA. A

combination of the anchoring and tagging enzymes yields a 10 base pair tag which is unique to a gene.

[0074] Using this approach, sequence tags for each gene were represented by the 3'-most NlaIII site followed by a unique 10 bp sequence. In instances where tags could not be assigned to known genes, a SAGE library cDNA was PCR amplified using the SAGE tag and a commonly used M13 forward primer (GTTTTCCTCAGTCACGAC, SEQ ID NO:18). PCR products were subsequently cloned into the pCR2.1 vector (Invitrogen) and sequenced using standard techniques. Identification of genes was based on the homology of the sequence of PCR products to known sequences in GenBank® (www.ncbi.nlm.nih.gov/genbank).

[0075] A BLAST® alignment (www.ncbi.nlm.nih.gov/blast) of nucleotide sequences to their mouse and/or rat counterparts was performed to identify the gene from which the tag was derived. Of the sixteen most abundant tags identified in this analysis (Table 2), the gene for all but one tag was identified. Of these fifteen identified genes, five were mitochondrial in origin and three were nuclear repetitive elements. Occurrence of multiple copies of these genes in each cell was the likely cause of their abundance in the SAGE output. Such sequences were not considered for further evaluation.

TABLE 2

Abundance	Tag	Gene	SEQ ID NO:	Identified
38	CATGGAAGCAGAAT	Alu Repeat	19	J00052
33	CATGCAGGAGCTTC	Mito COX I	20	PCR
27	CATGGGGGAGCGTT	Ribosomal Protein S21	21	PCR
27	CATGGTACTGACAC	Mito COX III	22	PCR
20	CATGGCCTCCAAGG	GAPDH	23	X52123
20	CATGATAATACGTA	Mito ATPase 6	24	M14311
19	CATGCCTTTAATCC	B-1 Repeat	25	PCR
18	CATGAATCGGAGGC	Mito Cytochrome B	26	J01436
18	CATGAGGCAGACAG	EF-1	27	D00522
18	CATGGCGGCAGACG	Galectin (L-14)	28	M96676
16	CATGGTGGCTCACA	Alu Repeat	29	J00056
15	CATGTTGGCTGCCG	Ferritin Heavy Chain	30	M99692
14	CATGCCCTGTGCCG	No Match	31	
13	CATGAGAGCGAAGT	Ribosomal Protein L41	32	X82550
13	CATGAGGAGGCCTA	Mitochondrial NADH Dehydrogenase	33	PCR
12	CATGCCCTGAGTCC	β -Actin	34	AF014363

[0076] Using this approach, promoters of four genes were identified as being the most active in CHO-K1 cells. These promoters were: β -actin, ribosomal protein S21 (rpS21), elongation factor 1 (EF-1), and glyceraldehyde 3-phosphate dehydrogenase (GAPDH). The high levels of these mRNAs in CHO-K1 cells could either be due to the promoter activity of their respective promoters or due to innate stability of the mRNAs. Although SAGE analysis provides a quantification of overall steady state levels for the mRNAs for genes, it does not distinguish between promoter activity of the gene and mRNA stability as the basis of the high expression of the mRNA. Thus, in order to distinguish between the two possibilities, half-life of mRNAs

were measured. Briefly, expression of candidate genes was assessed by Northern blot analysis of CHO-K1 cells in spinner cultures at varying points following treatment of cells with actinomycin D.

[0077] Initially, the rpS21, GAPDH and EF-1 genes were analyzed and were all found to have relatively stable mRNAs with half-lives greater than 8 hours. These results suggested that the greater abundance of these mRNAs resulted from greater stability of the mRNAs and not necessarily greater activities of the respective promoters.

[0078] The half-life of galectin, ferritin, and β -actin mRNAs was also measured by Northern blot analysis, as described above, at 0, 4, 8, 10, and 15 hours following treatment of cells with actinomycin D. A representative Northern blot is shown in Figure 6A. The relative mRNA levels are represented graphically in Figure 6B. These data show that although both galectin and ferritin had half-lives of greater than 8 hours, the β -actin mRNA turned over more rapidly with a half-life of approximately 6 hours. Thus, the relative contribution of promoter strength to overall steady state mRNA levels was greater for β -actin than the other candidates in CHO-K1 cells. Accordingly, under these conditions, the β -actin promoter can be characterized as a strong promoter.

Example 2: Isolation and Characterization of the Hamster β -Actin and rpS21 Promoters

[0079] In light of the results described in Example 1, the candidate with the greatest abundance (rpS21) and the one with the most rapid mRNA

turnover (β -actin) were selected for further study. A λ FIX II CHO-K1 genomic library (Stratagene, LaJolla, CA) was screened to isolate genomic DNAs for hamster β -actin and rpS21 promoters.

[0080] In order to isolate β -actin and rpS21 genomic clones, the E. coli bacterial strains, XL1-Blue MRA (P2) were grown in LB medium containing 10 mM magnesium sulfate and 0.2% maltose. The bacterial cells were pelleted and resuspended in 10 mM magnesium sulfate at an absorbance reading of 0.5 at 600 nm. Approximately one million phage from the library were incubated with the bacterial cells for 15 minutes at 37°C. Molten agarose was added to the phage/bacteria mixture and the bacteria were overlayed on agar-containing BioAssay plates (Nunc, Rochester, NY). Following the hardening of the top agarose, the plates were inverted and grown at 30°C overnight. Plates were subsequently chilled and overlayed twice with Genescreen Plus™ nylon filters (Perkin Elmer Life Sciences, Wellesley, MA). The nylon filters were denatured for 2 minutes in 0.1 M sodium hydroxide with 1.5 M sodium chloride and subsequently neutralized. Filters were UV cross-linked and probed.

[0081] A probe used for isolation of the hamster β -actin promoter was derived by random PCR from the 5' end of the β -actin gene (nt 238-381 of GenBank® Accession No. U20114). A probe used for the isolation of hamster rpS21 promoter was derived by PCR using primers set forth in SEQ ID NOs:12 and 13. Hybridizing phage for both β -actin and rpS21 promoters were purified using standard techniques. The DNA from the phage isolated

from the phage lysates was purified by sequential extractions with chloroform, phenol, phenol/chloroform (1:1), and lastly, chloroform.

[0082] For isolation of hamster β -actin gene promoters, following ethanol precipitation, DNA was digested with restriction enzymes that had sites in the 5' portion of the β -actin hamster gene and subjected to Southern blotting using the same probe that was used to screen the genomic library.

[0083] Using this approach, an *AvrII* fragment of approximately 7 kb and a *Sall* fragment of approximately 5.5 kb were generated, both of which hybridized to the probe. These were subsequently cloned into pBluescript II KS plasmid (Stratagene). The 7 kb *AvrII* fragment has the ATCC Reference No. PTA-5309, deposited July 3, 2003 with the American Tissue Culture Collection, P.O. Box 1549, Manassas, VA 20108, U.S.A.

[0084] Plasmids containing *AvrII* and *Sall* fragments were digested with *SfiI* to remove the 3' end of the fragments which contained a portion of the open reading frame of the β -actin gene. These fragments were then cloned into the pDsRED-1 plasmid (Clontech) to create the constructs termed pDsRED-Avr (6.5kb) and pDsRED-Avr (5.1kb). In order to generate a construct containing all of intron 1 of the β -actin gene, PCR was performed using the following primers:

Forward: AGGCCAGCTTGGGACCAAGACAGAA (SEQ ID NO:35)

Reverse: CGCGGATCCGGCGAACTATATCAGGGC (SEQ ID NO:36).

[0085] The PCR fragment generated two products: a predicted product of approximately 7 kb and a smaller unexpected 3 kb product. Both

of these PCR products were cloned into the pDsRED-1 plasmid (Clontech) to generate the constructs pDsRED-Avr(1)-7 and pDsRED-Avr(1)-3.

[0086] Each of the fragments of the β -actin hamster promoter that were cloned into the pDsRED-1 plasmid (Clontech) were transfected into CHO-K1 cells. The relative promoter strengths of each of the hamster β -actin promoter fragments were measured using FACS as described above. The results of the activity assays are summarized below.

[0087] Avr(1)-3 fragment of β -actin promoter which spans from nt -1970 to nt +1037 exhibited the highest promoter activity. The Avr(1)-7 fragment which spans from nt -6000 to nt +1037 exhibited an activity that was 47% of the activity exhibited by Avr(1)-3. The Avr(6.5 Kb), Sal(5.1Kb), Actin(3 kb), and Actin-P(2.8 kb) fragments exhibited only 2%, 2%, 2%, and 0% promoter activity, respectively, as compared to the Avr(1)-3 fragment.

[0088] The Avr(1)-3 fragment was subsequently sequenced, and the sequence is set forth in SEQ ID NO:1. Additionally, the region 660 nt upstream of the 5' of Avr(1) 3 was also sequenced. This longer sequence from nt -2622 to nt +1037 is set forth in SEQ ID NO:7.

[0089] For isolation of the *rpS21* promoter, following isolation of DNA from the hybridizing phage, the DNA was amplified by PCR using the following primers:

Forward: AGCTCTAATACGACTCACTATAGGGC (SEQ ID NO:40)

Reverse: CTCTAGGCCAGCGGAGCGCAG (SEQ ID NO:41).

The PCR product was cloned into the vector PCR2.1 (Invitrogen) and subsequently sequenced. The nucleotide sequence of the hamster rpS21 promoter is set forth in SEQ ID NO:39. The promoter was excised using EcoRI sites flanking the cloning sites and cloned into the pDsRED1-1 vector (Clontech). The 2 kb hamster rpS21 promoter sequence has ATCC Reference No. _____, deposited _____, 200____, with the American Tissue Culture Collection, P.O. Box 1549, Manassas, VA 20108, U.S.A.

Example 3: Functional Comparison of the Hamster β -Actin and CMV Promoters

[0090] The promoter activity of Avr(1)-3 was compared to that of the CMV immediate early promoter (Invitrogen) and the human EF-1 promoter (Invivogen).

[0091] CHO-K1 cells were transiently transfected with either pDsRED-1 plasmid containing either Avr(1)-3, the CMV immediate early promoter upstream, or the human EF-1 promoter, each operably linked to the RFP gene. Expression of RFP was assessed by FACS 48 hours post-transfection.

[0092] As shown in Figure 7A, in cells transfected with Avr(1)-3, the β -actin promoter sequence (SEQ ID NO:1) showed a higher level of RFP expression as compared to either the CMV or EF-1 promoters. In particular, expression was approximately two-fold higher with Avr(1)-3 than with the CMV promoter.

[0093] In order to determine whether this observed expression profile is sustainable in stable transfectants, transfected CHO-K1 cells were selected for two weeks with G418TM. Expression of RFP in the surviving pools of cells was then assessed. As depicted in Figure 7B, similarly to transiently transfected cells, the highest RFP expression was observed in cells transfected with Avr(1)-3, the β -actin promoter sequence set forth in SEQ ID NO:1. Example 4: Activity of the Hamster β -Actin Promoter in BHK-21 and HEK293 cells

[0094] The activity of the hamster β -actin promoter was compared to that of CMV promoter in BHK-21 (ATCC No. CCL 10) and HEK293 (ATCC No. CRL-1573) cells using stable transfection assays as described in Example 3. As seen previously in CHO-K1 cells, expression of RFP in BHK-21 cells was significantly higher when using the β -actin promoter instead of the CMV promoter (Table 3). In HEK293 cells, the hamster β -actin promoter resulted in expression of RFP at levels roughly equivalent to those of the CMV promoter.

TABLE 3

Cell line	CMV promoter	β -Actin promoter
BHK-21	8.3 ± 0.4	121 ± 99.8
HEK293	139 ± 9.9	102 ± 8.3

Example 5: Rat and Mouse β -Actin Promoters

[0095] Publicly available databases of nucleotide sequences were searched using default settings for potential homologs of the hamster β -actin promoter sequence set forth in SEQ ID NO:1.

[0096] The 5' portion of a β -actin hamster gene (GenBank® Accession No. U21104; SEQ ID NO:4) exhibits 98% identity to the 3' portion of the hamster β -actin promoter sequence. This homology, however, is only 40% over the entire length of the hamster β -actin promoter sequence set forth in SEQ ID NO:1. No promoter activity is known for this portion.

[0097] Previously known β -actin promoters: human (GenBank® Accession No. gi28337A) and chicken (GenBank® Accession No. gi2170437) were aligned with the hamster β -actin promoter for homology determination with the BLAST® program using default settings. The human and the chicken β -actin promoter sequences had only 10% and 1% identity, respectively, to the hamster β -actin promoter (SEQ ID NO:1).

[0098] A rat (*Rattus norvegicus*) genomic supercontig (GenBank® Accession No. NW_042778) was identified on chromosome 12 of the rat genome as containing a nucleotide sequence having a 67% identity over the entire length of SEQ ID NO:1.

[0099] Similarly, a contig (GenBank® Accession No. NT_039324) was identified on chromosome 5 of the mouse (*Mus musculus*) genome as having a 80% identity over the entire length of SEQ ID NO:1.

[0100] The sequence alignments of hamster β -actin promoter sequence (SEQ ID NO:1) with the hamster gene sequence, and β -actin promoters from human, chicken, rat and mouse are depicted in Figures 3, 4, 5, 1, and 2, respectively.

Example 6: Activities of the Rat and Mouse β -Actin Promoters

[0101] The rat and the mouse promoter sequences set forth in SEQ ID NOs:2 and 3, respectively, are cloned into the pDsRED-1 plasmid (Clontech). The CMV promoter is also cloned upstream of the RFP gene in the pDsRED-1 plasmid. These plasmids are transfected into CHO-K1 cell, or another cell line. Expression of the RFP is assessed by FACS 48 hours post-transfection.

[0102] Cells transfected with the rat or the mouse β -actin promoter are expected to show a higher RFP expression than the CMV promoter under similar conditions.

Example 7: Expression of Proteins Using Hamster β -Actin Promoter

[0103] To further evaluate activity of the hamster β -actin promoter, an expression system utilizing dihydrofolate reductase (DHFR) selection and methotrexate (MTX) amplification was used. The vector pGZ6 was derived from the pCLHAXSV2DHFR plasmid, so as to contain the 3 kb hamster β -actin promoter (SEQ ID NO:1) in addition to a DHFR gene under the control of the SV40 early promoter. The pCLHAXSV2DHFR plasmid has been previously described by Cole et al. (1993) *Biotechnology*, 11:1014-1024. Briefly, the metallothionine (MT) promoter in the pCLHAXSV2DHFR vector was replaced with the β -actin promoter to create the pGZ6 vector. cDNAs for two proteins of therapeutic interest, acid sphingomyelinase (ASM) and α -glucosidase (GAA) were operably linked to the hamster β -actin promoter. The ASM cDNA was obtained through the IMAGE™ consortium (GenBank® Accession No. AI587087). The cDNA for GAA was obtained from Dr. Martinuik at the New York University School of Medicine. The nucleotide sequences of the ASM and GAA cDNAs are set forth in SEQ ID NOs:37 and 38, respectively. Similarly, the two cDNAs were also cloned downstream of the CMV promoter in a vector containing the same DHFR expression cassette. The DHFR-deficient CHO-K1 cell line DXB11 was transfected in triplicate with both sets of expression vectors. After two weeks of selection in nucleotide-deficient media containing 20 nM MTX, a heterogeneous uncloned pools of cells were washed with PBS and transferred to serum-free media. Twenty four hours later, levels of ASM or GAA in the media were measured.

[0104] The results of one such experiment are demonstrated in Figures 8A and 8B. The levels of ASM generated from the hamster β -actin promoter in the stable pools were from 2 to 15 times greater than with the CMV promoter, and in the case of the GAA pools, 2 to 5 times greater.

[0105] The stable pools were further used to evaluate the ability of the β -actin promoter to sustain long-term protein expression. Typically, for industrial production of proteins, high expression is achieved by selecting cells with a higher gene copy number through a process that involves increasing the number of selection steps and/or concentration of MTX. In order to determine whether a higher expression could be achieved via this strategy with the β -actin promoter (SEQ ID NO:1), the ASM pools initially selected at 20 nM MTX were amplified by selection for two weeks at ten-fold higher levels of MTX (200 nM). As summarized in Table 4, two of the three tested β -actin pools showed 2 to 3-fold greater levels of ASM after amplification relative to the starting 20 nM pools. In contrast, only one of the CMV pools tested showed higher levels than the 20 nM pool, from which it was derived. Among the six ASM pools generated with either of the two promoters, the highest expressing β -actin pool generated six times the amount of ASM obtained with the highest expressing pool generated with CMV promoter. This demonstrates that, at least under the conditions tested, the hamster β -actin promoter is superior to the CMV promoter.

TABLE 4

Pool	Expression of ASM at 20 nM MTX	Expression of ASM at 200 nM MTX
CMV-ASM Pool A	4.3	8.2
CMV-ASM Pool B	16.9	9.5
CMV-ASM Pool C	3.6	3.7
β -actin-ASM Pool A	33.5	100.0
β -actin-ASM Pool B	59.3	27.9
β -actin-ASM Pool C	45.6	90.5

[0106] In a separate experiment, the hamster β -actin promoter was used for expressing tissue plasminogen activator (tPA) protein, which is a thrombolytic agent used in patients for dissolving blood clots. CHO-DXB11 cells were transfected with a pGZ6-tPA expression vector in which the hamster β -actin promoter is operably linked to the tPA gene. Stable transfectants were selected by growth in nucleotide deficient medium containing 200 nM MTX. The resulting pool of uncloned cells was then subjected to 500 nM MTX to amplify transgene copy number. This pool of cells was removed from MTX, expanded and seeded on Cytopore™ 2 microcarriers in a 1 liter spinner culture. Cells were grown for 7 days in a serum containing medium. For the next 4 days, the serum was removed by daily 80% exchanges with serum free medium. Media harvests were then collected over 15 days and analyzed for tPA expression using a commercially available ELISA kit (TintElize® tPA kit, Biopool International, Inc., Ventura, CA). As depicted in Figure 9 of this experiment, the use of the hamster β -actin promoter resulted in tPA expression at a concentration of about 30

mg/L per day. This result compares favorably to recently published reports in which about 30-40 mg/L of tPA was produced after 4-8 days using other promoters (Senger et al. (2003) Biotechnology Progress 19: 1199-1209; Dowd et al. (2000) Biotechnology Progress 16:786-794).

Example 8: Production of Antibodies Using Hamster β -Actin Promoter

[0107] In order to produce an antibody to a TGF- β family member, nucleic acid encoding either an anti-TGF- β antibody light chain or an anti-TGF- β antibody heavy chain is cloned downstream of the hamster β -actin promoter in two separate pGZ6 expression vectors.

[0108] The DHFR-deficient CHO-K1 cell line DXB11 is transfected in with both expression vectors. After two weeks of selection in nucleotide-deficient media containing MTX, levels of anti-TGF- β antibody, including both the light chain and the heavy chain, are measured in the media.

Example 9: Expression of Proteins Using Hamster rpS21 Promoter

[0109] The hamster rpS21 promoter activity was compared to the hamster β -actin promoter activity for expression in CHO-DXB11 cells. CHO-DXB11 cells were transfected with expression vectors containing human α -glucosidase (rhGAA) operably linked to either the hamster rpS21 promoter of SEQ ID NO:39 (pGZ3IC-GAA) or hamster β -actin promoter of SEQ ID NO:1 (pGZ6IC-GAA). In both cases the rhGAA gene was linked to the gene encoding a cell surface marker (CD20) through an internal ribosome entry site

(IRES) sequence. After selection of cells with 0.2 μ M MTX in nucleotide deficient medium, the cells were labeled with a FITC-conjugated antibody to CD20 and sorted by FACS for high expressing clones. Selected cells were plated in 96-well plates and expanded for evaluation of rhGAA expression. 38 clones were analyzed for the hamster *rpS21* promoter, and 29 clones were analyzed for the hamster β -actin promoter. Table 5 shows the distribution of expression ranges in the resulting clones for both promoters.

TABLE 5

Vector	GAA Expression <2 pg/cell/hr	GAA Expression 2-5 pg/cell/hr	GAA Expression 5-8 pg/cell/hr	GAA Expression 8-10 pg/cell/hr
pGZ3IC-GAA	16%	50%	26%	8%
pGZ6IC-GAA	52%	34%	14%	0%

[0110] In a separate experiment, the hamster *rpS21* promoter was used for expressing ASM in CHO-DXB11 cells. The activity of the *rpS21* promoter was compared to activities of both β -actin and CMV promoters. CHO-DXB11 cells were transfected in triplicate and either selected directly at 200 nM MTX, or initially selected at 20 nM MTX and then amplified for two weeks at 200 nM MTX, as discussed in Example 7. Levels of ASM were measured in the media as described. ASM expression in untransfected cells was undetectable.

[0111] As summarized in Table 6, all three *rpS21* pools showed 2- to 3-fold greater levels of ASM after amplification relative to the starting 20 nM

pools, from which they were derived. Further, the levels of ASM generated were higher than the levels generated with the CMV promoter(Example 7).

TABLE 6

Pool	Expression of ASM nU/cell/24 hr (at 20 nM MTX)	Expression of ASM nU/cell/24 hr (at 200 nM MTX)
rpS21-ASM Pool A	12	34
rpS21-ASM Pool B	13	30
rpS21-ASM Pool C	16	41

[0112] The levels of ASM expression generated with selection of the pools directly at 200 nM MTX are summarized in Table 7.

TABLE 7

Pool	ASM Expression
CMV-ASM Pool A	38
CMV-ASM Pool B	193
CMV-ASM Pool C	44
β -actin-ASM Pool A	381
β -actin-ASM Pool B	125
β -actin-ASM Pool C	515
rpS21-ASM Pool A	342
rpS21-ASM Pool B	60
rpS21-ASM Pool C	51

[0113] The levels of ASM generated from the hamster rpS21 promoter at 200 nM MTX were on average about 1 to 2 times greater than that with the CMV promoter. The ASM levels generated from the β -actin promoter, on the other hand, were on average about 3 to 4 times greater than that with the CMV promoter. Thus, the rpS21 promoter was at least as active as the β -actin promoter when used for expressing GAA, however, it exhibited

lower activity than the β -actin promoter when used to express ASM. Both promoters, however, were more active than the CMV promoter.

[0114] The specification is most thoroughly understood in light of the teachings of the references cited within the specification which are hereby incorporated by reference. The embodiments within the specification provide an illustration of embodiments of the invention and should not be construed to limit the scope of the invention. The skilled artisan readily recognizes that many other embodiments are encompassed by the invention. All publications and patents cited and sequences identified by accession or database reference numbers in this disclosure are incorporated by reference in their entirety. To the extent the material incorporated by reference contradicts or is inconsistent with the present specification, the present specification will supercede any such material. The citation of any references herein is not an admission that such references are prior art to the present invention.

[0115] Unless otherwise indicated, all numbers expressing quantities of ingredients, cell culture, treatment conditions, and so forth used in the specification, including claims, are to be understood as being modified in all instances by the term "about." Accordingly, unless otherwise indicated to the contrary, the numerical parameters are approximations and may vary depending upon the desired properties sought to be obtained by the present invention. Unless otherwise indicated, the term "at least" preceding a series of elements is to be understood to refer to every element in the series. Those skilled in the art will recognize, or be able to ascertain using no more than

routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

CLAIMS

1. An isolated rodent β -actin promoter chosen from nucleotide sequences set forth in SEQ ID NOs:1, 2 and 3, or a variant thereof having promoter activity.
2. An isolated hamster β -actin promoter nucleotide sequence set forth in SEQ ID NO:1, or a variant thereof having promoter activity.
3. An isolated rat β -actin promoter nucleotide sequence set forth in SEQ ID NO:2, or a variant thereof having promoter activity.
4. An isolated mouse β -actin promoter nucleotide sequence set forth in SEQ ID NO:3, or a variant thereof having promoter activity.
5. An isolated nucleic acid comprising nucleotide sequence set forth in SEQ ID NO:1, or a variant thereof having promoter activity.
6. An isolated nucleic acid comprising nucleotide sequence set forth in SEQ ID NO:2, or a variant thereof having promoter activity.
7. A vector comprising the promoter of SEQ ID NO:1, or a variant thereof having promoter activity.
8. A vector comprising the promoter of SEQ ID NO:2, or a variant thereof having promoter activity.
9. A vector comprising the promoter of SEQ ID NO:3, or a variant thereof having promoter activity.
10. The vector of any one of claims 7-9, wherein the promoter is operably linked to a heterologous nucleic acid.

11. The vector of claim 10, wherein the heterologous nucleic acid encodes a therapeutic protein.
12. The vector of claim 11, wherein the therapeutic protein is chosen from acid sphingomyelinase, α -glucosidase, and tissue plasminogen activator.
13. A host cell transfected with a vector of any of claims 7-12.
14. The host cell of claim 13, wherein the cell is a CHO cell.
15. A method of producing a protein comprising:
 - (a) culturing a cell transfected with a vector comprising a hamster β -actin promoter, or a variant thereof, operably linked to a nucleic acid molecule encoding the protein; and
 - (b) recovering the protein.
16. The method of claim 15, wherein the protein is an antibody.
17. The method of claim 16, wherein the antibody binds a TGF- β family member.
18. The method of claim 15, wherein the protein is a therapeutic protein.
19. The method of claim 18, wherein the therapeutic protein is chosen from acid sphingomyelinase, α -glucosidase, and tissue plasminogen activator.
20. A transgenic animal comprising the promoter as in any one of claims 1-6.

21. The transgenic animal of claim 20, wherein the animal is a mammal.
22. An isolated rpS21 promoter having the nucleotide sequence set forth in SEQ ID NO:39, or a variant thereof having promoter activity.
23. A vector comprising the nucleotide sequence set forth in SEQ ID NO:39, or a variant thereof having promoter activity.
24. A vector of claim 23, wherein the nucleotide sequence is operably linked to a heterologous nucleic acid.
25. The vector of claim 24, wherein the heterologous nucleic acid encodes a therapeutic protein.
26. The vector of claim 25, wherein the therapeutic protein is α -glucosidase or acid sphingomyelinase.
27. A host cell transfected with a vector of any of claims 23-26.
28. The host cell of claim 27, wherein the cell is a CHO cell.
29. The method of producing a protein comprising:
 - (a) culturing a cell transfected with a vector comprising a hamster rpS21 promoter, or a variant thereof, operably linked to a nucleic acid molecule encoding the protein; and
 - (b) recovering the protein
30. The method of claim 29, wherein the protein is an antibody.
31. The method of claim 29, wherein the protein is a therapeutic protein.

32. The method of claim 31, wherein the therapeutic protein is α -glucosidase or acid sphingomyelinase.
33. A transgenic animal comprising the promoter as in claim 22.
34. The transgenic animal of claim 33, wherein the animal is a mammal.
35. An isolated β -actin promoter having a nucleotide sequence as deposited under ATCC Reference Number PTA-5309.
36. An isolated rpS21 promoter having a nucleotide sequence as deposited under ATCC Reference Number_____

[illegible]

FIG. 1B

FIG. 1B (CONT'D-1)

FIG. 1B (CONT'D-2)

[illegible]

FIG. 1B (CONT'D-3)

[illegible]

FIG. 1B (CONT'D-4)

FIG. 2A

[illegible]

FIG. 2B

FIG. 2B (CONT'D-1)

FIG. 2B (CONT'D-2)

FIG. 2B (CONT'D-3)

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FIG. 2B (CONT'D-4)

FIG. 3

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FIG. 3(CONT'D-1)

FIG. 3(CONT'D-2)

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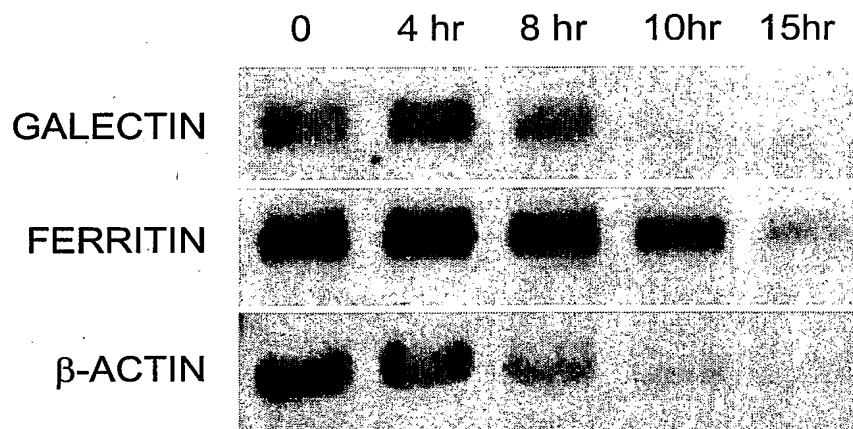
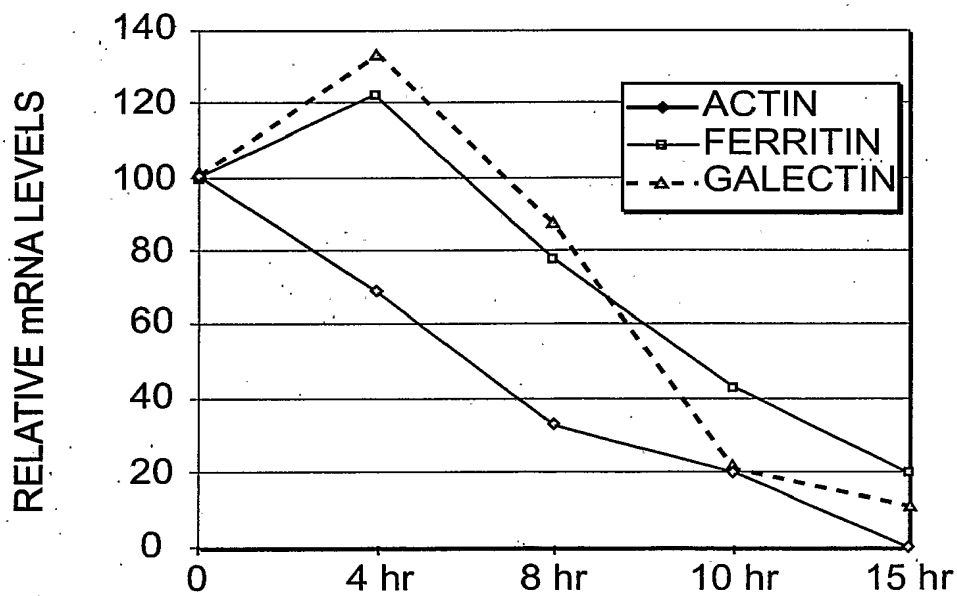
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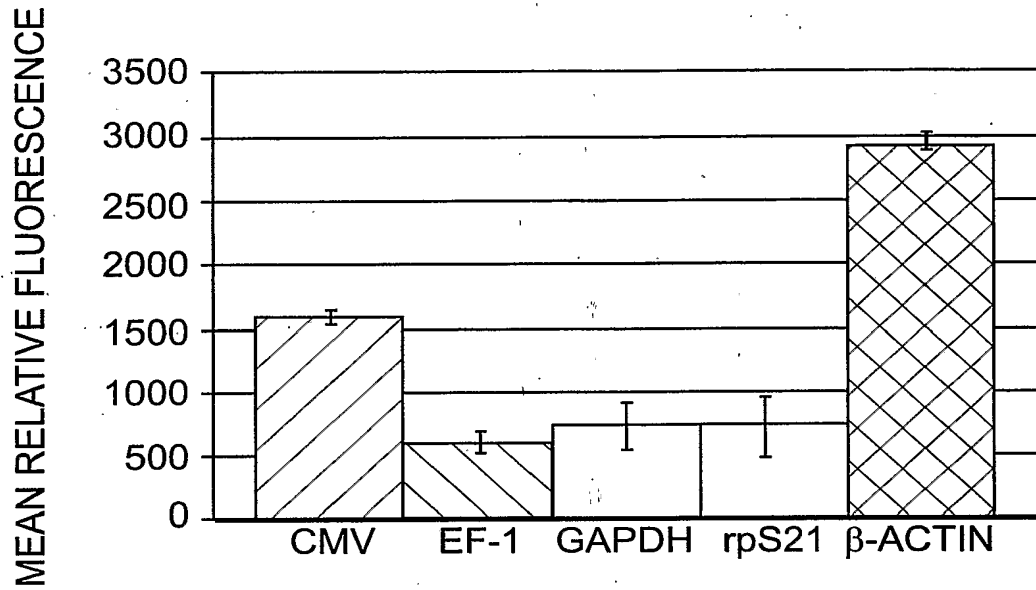
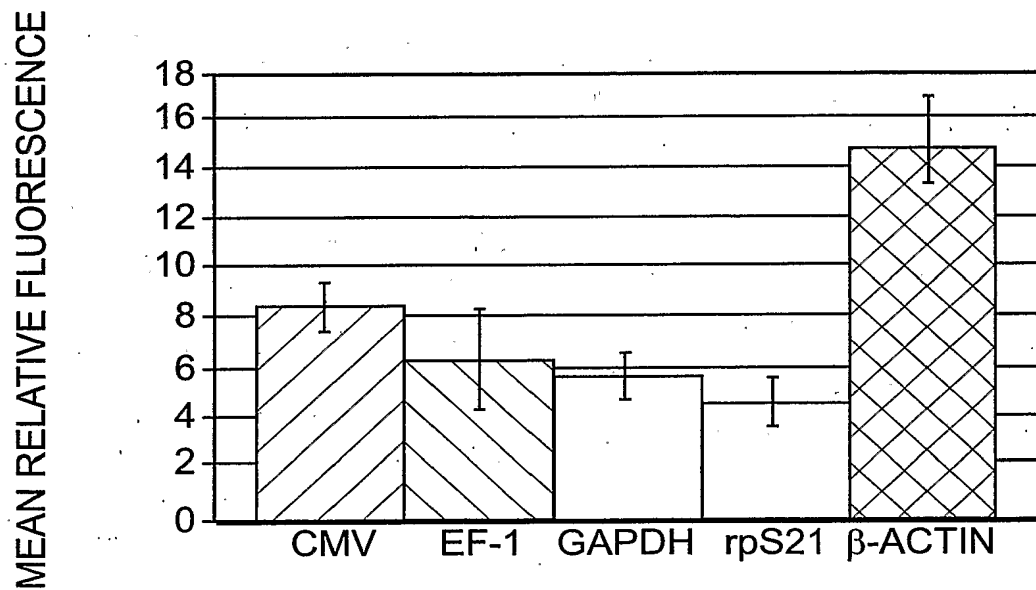
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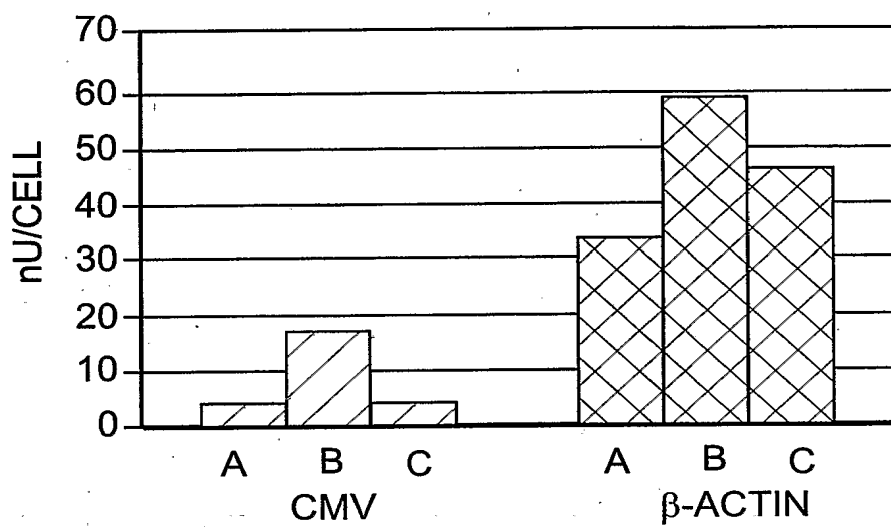
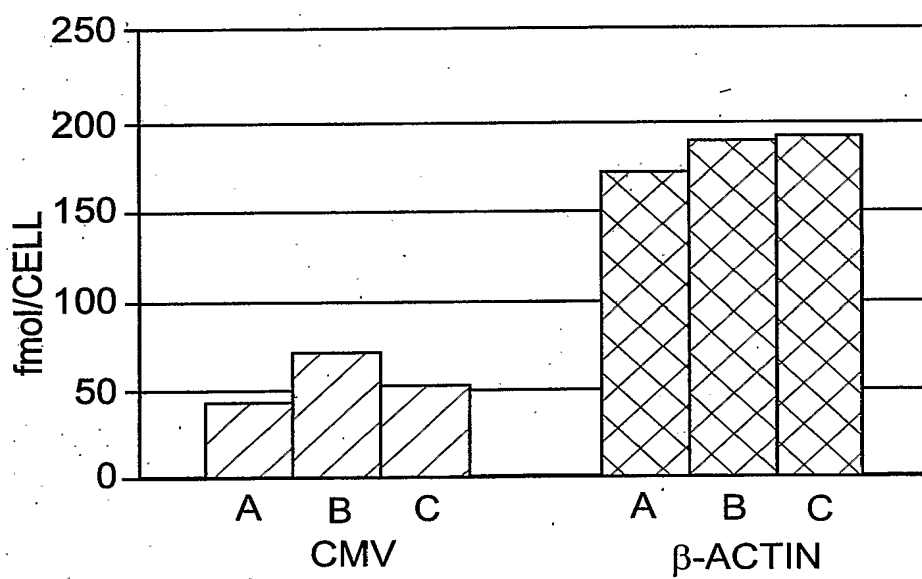
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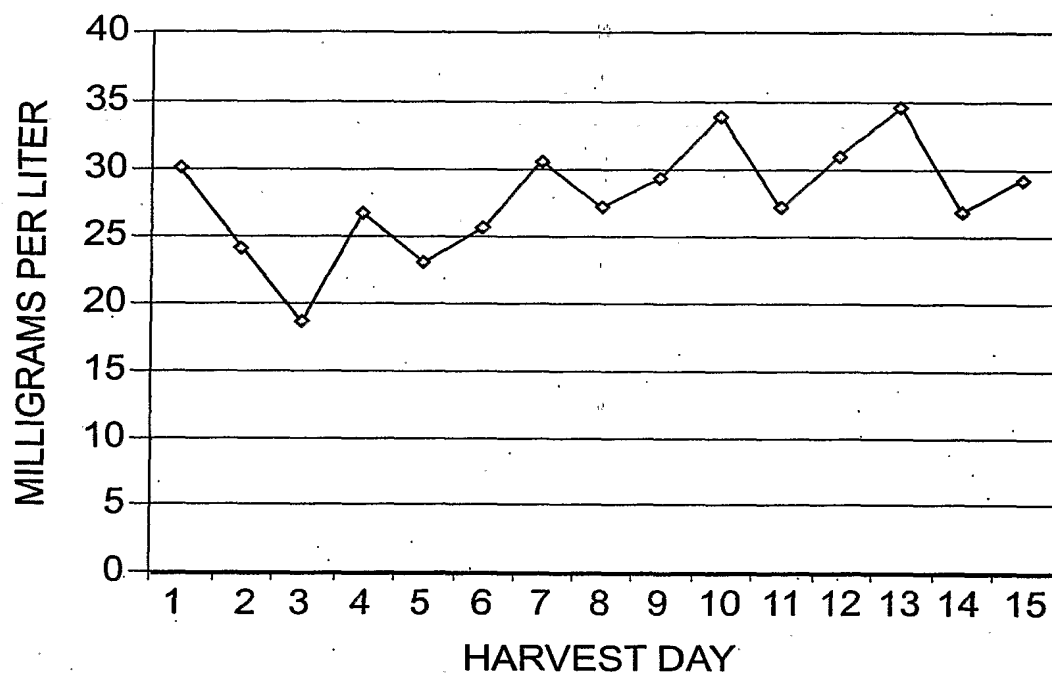
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FIG. 5

**FIG. 6A****FIG. 6B**

**FIG. 7A****FIG. 7B**

**FIG. 8A****FIG. 8B**

**FIG. 9**

SEQUENCE LISTING

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ZHANG, WEIQUN
GENZYME CORP.

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<213> Artificial Sequence

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<220>
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<400> 8
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<210> 9
<211> 19
<212> DNA
<213> Artificial Sequence

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<220>
<223> reverse primer for beta-actin

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<400> 9
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<210> 10
<211> 20
<212> DNA
<213> Artificial Sequence

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<220>
<223> forward primer for EF-1

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<400> 10
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<210> 11
<211> 17
<212> DNA
<213> Artificial Sequence

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<220>
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<210> 12
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<220>

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<210> 13
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<400> 13
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<210> 14
<211> 21
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<220>
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<400> 14
cgccagaact accaccagga c 21

<210> 15
<211> 20
<212> DNA
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<220>
<223> reverse primer for ferritin
<400> 15
ttcagagcca catcatcccg 20

<210> 16
<211> 21
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<220>
<223> forward primer for galectin
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tggtcgcaag caacctgaat c 21

<210> 17
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<220>
<223> reverse primer for galectin
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ttgaagtcac cgtctgccgc 20

<210> 18
<211> 17
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<220>

<223> forward M13 primer

<400> 18
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17

<210> 19
<211> 14
<212> DNA
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<220>
<223> alu repeat SAGE tag

<400> 19
catggaagca gaat

14

<210> 20
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<212> DNA
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<220>
<223> Mitochondrial COX I SAGE tag

<400> 20
catgcaggag cttc

14

<210> 21
<211> 14
<212> DNA
<213> Artificial Sequence

<220>
<223> Ribosomal Protein S21 SAGE tag

<400> 21
catgggggag cggt

14

<210> 22
<211> 14
<212> DNA
<213> Artificial Sequence

<220>
<223> Mitochondrial COX II SAGE tag

<400> 22
catggtactg acac

14

<210> 23
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<213> Artificial Sequence

<220>
<223> GAPDH SAGE tag

<400> 23
catggcctcc aagg

14

<210> 24
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<212> DNA
<213> Artificial Sequence

<220>

<223> Mitochondrial ATPase SAGE tag

<400> 24

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14

<210> 25

<211> 14

<212> DNA

<213> Artificial Sequence

<220>

<223> B-1 repeat SAGE tag

<400> 25

catgccttta atcc

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<210> 26

<211> 14

<212> DNA

<213> Artificial Sequence

<220>

<223> Mitochondrial Cytochrome B SAGE tag

<400> 26

catgaatcgg aggc

14

<210> 27

<211> 14

<212> DNA

<213> Artificial Sequence

<220>

<223> EF-1 SAGE tag

<400> 27

catgaggcag acag

14

<210> 28

<211> 14

<212> DNA

<213> Artificial Sequence

<220>

<223> Galectin SAGE tag

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<223> Alu repeat SAGE tag

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<210> 30

<211> 14

<212> DNA

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<223> Ferritin heavy chain SAGE tag
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 <223> Ribosomal protein L41 SAGE tag
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 <400> 33
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 <210> 34
 <211> 14
 <212> DNA
 <213> Artificial Sequence
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 <223> beta-actin SAGE tag
 <400> 34
 catgccctga gtcc 14
 <210> 35
 <211> 26
 <212> DNA
 <213> Artificial Sequence
 <220>
 <223> forward primer for amplifying beta-actin promoter containing
 intron 1
 <400> 35
 aggcccagct tgggaccaag acagaa 26
 <210> 36
 <211> 27
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> reverse primer for amplifying beta-actin promoter containing
 intron 1.

<400> 36
 cgcgatccg gcgaactata tcagggc

27

<210> 37
 <211> 1884
 <212> DNA
 <213> Artificial sequence

<220>
 <223> cDNA encoding acid-sphingomyelinase

<400> 37
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<210> 38

<211> 2859

<212> DNA

<213> Artificial Sequence

<220>

<223> cDNA encoding alpha-glucosidase.

<400> 38

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<210> 39
 <211> 1958
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Hamster rps21 promoter

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<210> 40
 <211> 26
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence: Synthetic primer

<400> 40
 agctctaata cgactcacta tagggc

26

<210> 41
 <211> 21
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence: Synthetic primer

<400> 41
ctctaggcca gcggagcgca g

21